U.S. PATENT DOCUMENTS

*EXAMINER INITIAL		DOCUMENT NO.	DATE	NAME	CLASS	SUB-CLASS	FILING DATE (if applicable)
NG	DA .	6,451,838 B1	9/17/2002	Moon et al			

FOREIGN PATENT DOCUMENTS

		DOCUMENT NO.	DATE	COUNTRY	CLASS	SUB-CLASS	TRANSLATION (yes/no)
NG	DB	01/90068	11/29/2001	wo			
NG	DC	03/015608	2/27/2003	WO			

OTHER REFERENCES (Including Author, Title, Date, Pertinent Pages, etc.)

NG	DE	Braud et al, "Potential Inhibitors of Angiogenesis. Part I: 3-(Imidazol-4(5)-ylmethylene)indoline-2-ones", Journal of Enzyme Inhibition and Medicinal Chemistry", Vol. 18, No. 3, June 2003, 243-252		
EYAMINED	/	Nyeemah Grazier/	DATE CONSIDERED	01/04/2007

EXAMINER / NYCEMAN Grazier/ DATE CONSIDERED 01/04/2007

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE FORM PTO-1449

NG

95, 138592n

LIST OF REFERENCES CITED BY APPLICANT

	LIST OF REFERENCES CITED BY APPLICANT			
ATTY. DOCKET:	SERIAL NO.:			
17543CON2(AP)	Not assigned			
APPLICANT:	TITLE: KINASE INHIBITORS FOR THE TREATMENT OF DISEASE			
Andrews et al				
FILING DATE:	GROUP:			
Submitted herewith	Not Assigned			
ILS PATENT DOCUMENTS				

*EXAMINER INITIAL		DOCUMENT NO.	DATE	NAME	CLASS	SUB-CLASS	FILING DATE (if applicable)
NG	AA_	4,966,849	10/30/1990	Vallee et al			
NG	AB	5,330,992	7/19/1994	Eissenstat et al			
NG	AC	5,217,999	6/8/1993	Levitzki et al			
NG	AD	5,302,606	4/12/1994	Spada et al			
NG	AE	5,792,783	8/11/1998	Tang et al			
NG	AF	5,834,504	11/10/1998	Tang et al			
NG	AG	5,883,113	3/16/1999	Tang et al			
NG	AH	5,883,116	3/16/1999	Tang et al			
NG	Al	5,886,020	3/23/1999	Tang et al			
NG	AJ	6,316,635	11/13/2001	Tang et al			
NG	AK	2002/0037878A1	3/28/2002	Moon et al			
NG	AL	2002/0035140A1	3/21/2002	Moon et al			*

		DOCUMENT NO.	DATE	COUNTRY	CLASS	SUB-CLASS	TRANSLATION (yes/no)
NG	AM	WO 94/10202	5/11/1994	PCT			
NG.	AN	WO 94/03427	2/17/1994	PCT			
NG	AO	WO 92/21660	12/10/1992	PCT			
NG	AP	WO 91/15495	10/17/1991	PCT			
- NG	AQ	WO 94/14808	7/7/1994	PCT			
NG	AR	WO 92/20642	11/26/1992	PCT			
NG	AS	WO 01/90103	11/29/2001	PCT			

OTHER REFERENCES (Including Author, Title, Date, Pertinent Pages, etc.) Plowman et al, "Receptor Tyrosine Kinases as Targets for Drug Intervention",1994, DN&P 7(6): 334-339 AT Bolen, "Nonreceptor tyrosine protein kinases", 1993, Oncogen 8: 2025-2031 AU Kendall et al, "Inhibition of vascular endothelial cell growth factor activity by an endogenously encoded soluble receptor", 1994, А۷ NG Proc. Natl'l Acad. Sci 90: 10705-10709 Kim et al, "Inhibition of vascular endothelial growth factor-induced angiogenesis suppresses tumor growth in vivo", Nature 362, AW NG Jellinek et al, "Inhibition of Receptor Binding by High-Affinity RNA Ligands to Vascular Endothelial Growth Factor", ΑX NG Biochemistry 33: 10450-10456 Takano et al, "Inhibition of Angiogenesis by a Novel Diaminoanthraquinone that Inhibits Protein Kinase C.", 1993, Mol. Bio. Cell ΑY NG 4: 2072, Page 358A AZ Kinsella et al, "Protein Kinase C Regulates Endothelial Cell Tube Formation on Basement Membrane Matrix, Matrigel", 1992, NG Experimental Cell Research, 199: 56-62 Wright et al, "Inibition of Angiogenesis In Vitro and In Ovo With an Inhibitor of Cellular Protein Kinases, MDL 27032", 1992, BA NG Journal of Cellular Phys. 152: 448-457 Mariani et al, "Inhibition of angiogenesis by FCE 26806, a potent tyrosine kinase inhibitor", 1994, Proc. Am. Assoc. Cancer Res. BB NG 35:2268; Page 381 Castro et al, "Quantitative Image Analysis of Laser-induced Choroidal Neovascularization in Rat", Exp. Eye Res. 2000; 71:523-55 BC NG BD Bundgaard et al, "Hydrolysis of N-(α-hydroxyalkyl)amide derivatives: implications for the design of N-acyloxyalkyl-type prodrugs", NG Int. J. of Pharmaceutics 22 (1984); 45-56 Bundgaard et al, ?Prodrugs as drug delivery systems, 43. O-Acyloxymethyl salicylamide N-Mannich bases as double prodrug forms BE NG for amines", Int. J. of Pharmaceutics 29 (1986); 19-28 Bundgaard et al, "A Novel Solution-Stable, Water-Soluble Prodrug Type for Drugs Containing a Hydroxyl or an NH-Acidic Group", BF NG J. Med. Chem. 32 (1989) 2503-2507 Bundgaard et al,"Prodrugs as drug delivery systems. XIX. Bioreversible derivatization of aromatic amines by formation of N-BG NG Mannich bases with succinimide", Chem. Abstracts 95, 138493f BH Bundgaard et al, "Hydrolysis of N-Mannich bases and its consequences for the biological testing of such agents", Chem. Abstracts

EXAMINER /Nyeemah Grazier/ DATE CONSIDERED 01/04/2007

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE FORM PTO-1449	Sheet2 of2_
LIST OF REF	ERENCES CITED BY APPLICANT
ATTY. DOCKET:	SERIAL NO.:
17543CON2(AP)	Not assigned
APPLICANT:	TITLE: KINASE INHIBITORS FOR THE TREATMENT OF DISEASE
Andrews et al	
FILING DATE:	GROUP:
Submitted herewith	Not Assigned

		OTHER REFERENCES (Including Author, Title, Date, Pertinent Pages, etc.)
NG	BI	Alminger et al, "(Pyridinylmethyl)sulfinylbenzimidazole derivatives as antiulcer agents, their preparation and formulations containing them", Chem. Abstracts 110, 57664p
NG	BJ	Buur et al, "Prodrugs of cimetidine with increased lipophilicity; N-acyloxymethyl and N-alkoxycarbonyl derivatives", Chem. Abstracts 115, 64029s
NG	BK	Hansen et al, "Carbamate ester prodrugs of dopaminergic compounds: synthesis, stability, and bioconversion", Chem Abstracts 115, 189582y
NG	BL	Bundgaard et al, "Phenyl carbamates of amino acids as prodrugs forms for protecting phenols against first-pass metabolism", Chem. Abstracts 117, 14347q
NG	ВМ	Jensen et al, N-Substituted (aminomethyl)benzoate 21-esters of corticosteroids as water-soluble, solution-stable and biolabile prodrugs", Chem. Abstracts 117, 55790x
NG	BN	Thomsen et al, "Evaluation of phenyl carbamates of ethyl diamines as cyclization-activated prodrug forms for protecting phenols against first-pass metabolism", Chem Abstracts 123, 17593b

/Nyeemah Grazier/	DATE CONSIDERED 01/04/2007					
*EXAMINER: Initial if reference considered, whether or not citation is in con	formance with MPEP 609; Draw line through citation if not in conformance and not					
considered. Include copy of this form with next communication to applicant.						